

*Free Home Sample Collection 9999 778 778

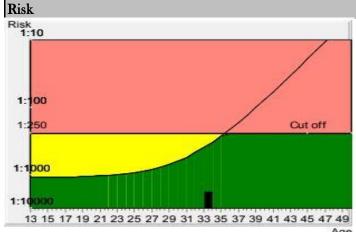


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Date of Report 22/10/19
PRISCA 5.0.2.37

					IMSCA	5.0.2.57	
Patient Data							
Name		Mrs Nidhi Gupta		Patient ID		011910200170	
Birthday		10/4/1986		Sample ID		10607591	
Age at delivery		33.5		Sample Date		21/10/19	
Gestational age		12+0					
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	72	Diabetes		no	Pregnancies		
Smoker	no	Origin		Asian			

Biochemical Data			Ultrasound Data		
Parameter	rameter Value		Gestational age	11+5	
PAPP-A	$1.68~\mathrm{mIU/ml}$	0.57	Method	CRL (<>Robinson)	
fb-hCG	30.3 ng/ml	0.65	Scan Date	19/10/19	
Risks at sampling date			Crown Rump Length (mm)	49	
Age Risk		1:361	Nuchal translucency MoM	0.53	
Biochemical Trisomy 21 Risk		1:1507	Nasal Bone	present	
Combined Trisomy 21 Ris	sk	1:8151	Sonographer	DR. PRAKASH LALCHANDANI	
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MD	



Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 8151 women with the same data, there is one woman with a trisomy 21 pregnancy and 8150 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

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